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(54) Emollient durability enhancing siloxanes.

⑤ A method of enhancing the durability of an emollient material on a substrate by forming a mixture of an emollient material and an effective amount of an organosilicon compound, and applying the mixture to the substrate to be treated. The organosilicon compound is either an aminofunctional, amidofunctional, or carboxyfunctional, polysiloxane. A skin conditioning composition of enhanced durability is also disclosed which is a mixture of an emollient material and an effective amount of an organosilicon compound which is either an aminofunctional, amidofunctional, or carboxyfunctional, polysiloxane.

This invention relates to the enhancement of the durability of emollient materials with certain organosilicon compounds. More particularly, the durability of mineral oil, for example, on human skin is enhanced by mixing the emollient with one of aminofunctional, amidofunctional, or carboxyfunctional, polysiloxanes.

This invention relates to a method of enhancing the durability of an emollient material on a substrate by forming a mixture of an emollient material and an effective amount of an organosilicon compound, and applying the mixture to the substrate to be treated. The organosilicon compound is either an aminofunctional, amidofunctional, or carboxyfunctional, polysiloxane.

Preferred embodiments of the present invention include an aminofunctional polysiloxane having the formula $Me_3SiO(Me_2SiO)_x(MeRSiO)_ySiMe_3$ in which Me is methyl; R is the functional group $-(C_4H_8)NH-(CH_2)_2NH_2$; x is an integer from fifty to one thousand; and y is an integer from one to fifty preferably one to twenty. The amidofunctional polysiloxane preferably has the formula $Me_3SiO(Me_2SiO)_x(MeRSiO)_ySiMe_3$ in which Me is methyl; R is the functional group $-(C_4H_8)NH(CH_2)_2NHCOCH_3$; x is an integer from fifty to one thousand preferably fifty to one hundred; and y is an integer from one to fifty preferably one to ten. Preferably, the carboxyfunctional polysiloxane compound has the formula $Me_3SiO(Me_2SiO)_x(MeRSiO)_ySiMe_3$ in which Me is methyl; R is the functional group $-(C_3H_6)COOH$; x is an integer from fifty to one thousand preferably one hundred to three hundred; and y is an integer from one to fifty.

The emollient and the polysiloxane may be present in the mixture in the ratio of four to one, more preferably two to one. The mixture of the emollient and the polysiloxane may, if desired, include a volatile cyclic siloxane solvent, and the solvent can be a low viscosity polydimethylcyclosiloxane fluid, for example, which is a mixture of cyclic tetramers and pentamers and having a viscosity of about 2.5 centistokes measured at 25° C. The mixture should contain about five to ten percent by weight of the emollient and the polysiloxane in the volatile cyclic siloxane solvent. The emollient may be a material such as mineral oil, mink oil, lanolin oil, and petrolatum.

The invention also relates to a skin conditioning composition of enhanced durability which is a mixture of an emollient material and an effective amount of an organosilicon compound. The organosilicon compound, as noted above, is either an aminofunctional, amidofunctional, or carboxyfunctional, polysiloxane.

In accordance with the present invention, there is provided a concept which relates to the enhancement of the durability of emollient materials with certain organosilicon compounds. The durability on human skin is enhanced by mixing an emollient with either one of an aminofunctional, amidofunctional, or carboxyfunctional, polysiloxanes. These aminofunctional, amidofunctional, and carboxyfunctional, organosilicon compounds are well known in the prior art, and such compounds as well as methods for preparing these compounds can be found in U.S. Patent No. 4,477,514, which shows carboxyfunctional siloxanes; U.S. Patent No. 4,559,227, which shows aminofunctional siloxanes; and U.S. Patent No. 4,848,981, which shows the amidofunctional siloxanes, of the present invention. A brief description of these functional siloxanes is set forth hereinbelow.

The amine functional siloxane polymer has the formula

 R_{3-z} 'Q_zSiO[R₂'SiO]_x[R'QSiO]_ySiQ_zR_{3-z}'

wherein R' denotes an alkyl group of 1 to 4 carbons or a phenyl group, with the proviso that at least 50 percent of the total R' groups are methyl; Q denotes an amine functional substituent of the formula —R"Z, wherein R" is a divalent alkylene radical of 3 to 6 carbon atoms or a radical of the formulation —CH₂CH₂CH₂-CHOHCH₂— and Z is a monovalent radical selected from the group consisting of —NR₂"", —NR""(CH₂)_nNR₂""; and

wherein R''' denotes hydrogen or an alkyl group of 1 to 4 carbons, R'''' denotes an alkyl group of 1 to 4 carbons and n is a positive integer from 2 to 6; z has a value of 0 or 1; x has an average value of 25 to 3000; y has an average value of 0 to 100 when z is 1, and y has an average value of 1 to 100 when z is 0.

The amidofunctional polysiloxane is a triorganosiloxane-endblocked polydiorganosiloxane having an average of 50 to 1000 siloxane units per molecule with an average of 1 to 50 of the siloxane units per molecule being amide-containing siloxane units. The amide-containing siloxane units bear a substituent of

the formula

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wherein n is 0 or 1, R' denotes an alkylene radical of 3 to 6 carbon atoms, and R" denotes a hydrogen radical or an alkyl radical of 1 to 6 carbon atoms, X denotes an acyl radical of the formula

X' denotes a hydrogen radical or X, and R''' denotes an alkyl radical of 1 to 4 carbon atoms and substantially all other organic substituents in the polydiorganosiloxane being methyl groups.

The amidofunctional silicone component in accordance with this invention consists essentially of a triorganosiloxane-endblocked polydiorganosiloxane which contains amidoalkyl substituents. Triorganosiloxane-endblocked polydiorganosiloxanes (amidofunctional silicone) consist essentially of terminal triorganosiloxane units of the formula $R_3 SiO_{1/2}$ and backbone diorganosiloxane units of the formula $R_2 SiO_{2/2}$. Trace amounts of other siloxane units in amidofunctional silicone, such as $SiO_{4/2}$ and $RSiO_{3/2}$, which are normally present as impurities in commercial polydiorganosiloxanes may be present. Preferably there are no $SiO_{4/2}$ units or $RSiO_{3/2}$ units in the amidofunctional silicones.

The R radicals of the above siloxane units are substantially either amide-containing radicals of the formula

or methyl radicals. Minor amounts of other organic substituents which are normally present as impurities in commercial polydiorganosiloxanes may be present. It should be understood, for example, that the amidofunctional silicones of this invention are often prepared by acylation of corresponding aminofunctional silicones. Consequently, the amidofunctional silicones may also contain residual aminofunctional siloxane units. For example, siloxane unites such as H₂NCH₂CH₂NHCH₂CH(CH₃)CH₂SiO_{2/2} or H₂NCH₂CH₂CH₂SiO_{2/2} may also be present in the amidofunctional silicones useful in this invention. However, for the purposes of this invention it is preferred to employ silicone oils that do not contain significant levels (more than 25 percent of the number of amidofunctional substituents) of the unmodified aminofunctional siloxane units.

In the formula for the amide-containing radicals, R' denotes an alkylene radical of 3 to 6 carbon atoms, such as -CH₂CH₂CH₂-, -CH₂CH₂CH₂-, -CH₂CH₂CH₂-, -CH₂CH₂CH₂-, -CH₂CH₂CH₂-, and -CH₂CH₂CH₂-(CH₂CH₃)CH₂-. Amidofunctional silicones wherein the silicon bonded, amide-containing radicals have a trimethylene radical or an alkylated trimethylene radical, such as -CH₂CH(CH₃)CH₂-, as the R' radical are preferred because of ease of synthesis and availability.

R" denotes a hydrogen radical, which is a preferred R" radical, or an alkyl radical of 1 to 6 carbon atoms, such as methyl, ethyl, propyl, butyl, and isobutyl.

In the formula for the amide-containing radicals, n has a value of 0 or 1, so that the radical may contain one or two nitrogen atoms. X denotes an acyl radical of the formula

and X' denotes a hydrogen radical or X. In the acyl radical, R'" denotes an alkyl radical of 1 to 4 carbon atoms such as methyl, ethyl, propyl, isopropyl, or butyl.

In accordance with the above, triorganosiloxane-endblocked polydiorganosiloxanes preferred for use in the method of this invention consists essentially of siloxane units selected from the following:

 $(CH_3)_3SiO_{1/2}$, and $(CH_3)_2SiO_{2/2}$ where R', R", and R"' have the same meanings as described above. It should be understood that any of the siloxane units having non-acylated nitrogen atoms can also be present in their salt form. It is well known that the salt form occurs when such polymers are neutralized by acids such as mineral acids or carboxylic acids.

The silicone polymers of this invention may contain amide-containing siloxane units of the formula

wherein R', R", and R" have the same meanings as described above. These amide-containing units have a ratio of acyl groups to nitrogen atoms of about 0.5.

The carboxyfunctional silicones of the invention have the formula

QMe₂SiO(Me₂SiO)_x(MeRSiO)_ySiMe₂Q

wherein Me is a methyl radical, R is a carboxyfunctional radical, said carboxyfunctional radical being selected from the group consisting of carboxyalkyl radicals and carboxythioalkyl radicals, Q is selected from the group consisting of R, Me and OH groups, x has a value of 1 to 1000, and y has a value of 1 to 100.

As referred to herein, a carboxyfunctional radical is a monovalent radical which contains the —COOH radical, and is attached to a silicon atom of the main molecular chain by a divalent linking group. Direct attachment to the silicon atom is through a silicon to carbon bond.

Divalent linking groups contemplated for use in the present invention are either alkylene groups containing from 2 to 9 carbon atoms, or thioalkylene groups, containing 2 to 8 carbon atoms and one sulfur atom present as a thioether group.

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Those carboxyfunctional radicals wherein the divalent linking group is an alkylene group are referred to herein as carboxyalkyl radicals; those carboxyfunctional radicals wherein the divalent linking group is a thioalkylene group are referred to herein as carboxythioalkyl radicals.

Specific examples of carboxyalkyl radicals include, but are not limited to $-CH_2CH_2COOH$, $-CH_2CH_3COOH$, $-CH_2CH_3COOH$, $-CH_2CH_3COOH$, $-CH_3COOH$, and the like. The $-CH_3COOH$ radical is a preferred carboxyalkyl radical for the practice of the present invention.

Specific examples of carboxythioalkyl radicals include, but are not limited to $-CH_2CH_2SCOOH$, $-(CH_2)_3SCOOH$, $-CH_2CH(CH_3)SCH_2COOH$, $-CH_2CH_2SCH_2COOH$, $-CH_2CH(C_2H_5)SCH_2COOH$, and the like. The $-CH_2CH_2SCH_2COOH$ radical is a preferred carboxythioalkyl radical for the practice of the present invention.

Examples of emollients and moisturizers which may be used in this invention include straight, branched or cyclic hydroxy compounds such as alcohols containing 1 to 30 carbon atoms; straight, branched, or cyclic carboxylic acids containing 1 to 31 carbon atoms; acid esters containing C₁ to C₃₀ carboxylic acids esterfied with C₁ to C₃₀ alcohols; alcohol ethers containing 1 to 30 carbon atoms; alkanes of the formula H—(CH₂)n—H, wherein n is 5 to 30; and siloxanes. Examples of such functional materials include 2-ethylhexyl oxystearate; arachidyl propionate; 2-ethylhexyl adipate; isopropyl myristate; ethanol; stearyl alcohol; propylene glycol; propionic acid; stearic acid; polyoxypropylene cetyl alcohol; polyoxypropylene lanolin alcohol; Carbowax® 300; petroleum jelly; mineral oil; aliphatic hydrocarbons such as mineral spirits; lanolin and lanolin derivatives such as acetylated lanolin and isopropyl lanolate; hexamethyldisiloxane; cyclic polydimethylsiloxane; linear polydimethylsiloxane; polypheylmethylsiloxane; and poly dimethyl/trimethylsiloxane. Other phenyl, ethyl and vinyl substituted polysilanes may also be included in the products of this invention.

In order to illustrate the durability enhancement of the siloxanes of the present invention, durability enhancement data was collected for a variety of emollients. A soap washing test procedure was used in order to measure the influence of the silicones on the durability of the various emollients. The emollients considered were mineral oil, mink oil, lanolin oil, and petrolatum. The test procedure was used to measure silicone substantivity on human skin. Specifically, the method was based on Attenuated Total Reflectance/Fourier Transform Infrared Spectrophotometric (ATR/FTIR) analysis, in which prism skin studies were conducted and analyzed based on the reflection of energy at the interface. Instrumentation included a NICOLET Model 20DX FTIR system, and a HARRICK Scientific Skin Analyzer. The ATR studies involved contact of the skin sample and prism. A hydration procedure was employed in order to increase the softness and flexibility of the skin surface which resulted in a less variable contact between the skin and prism. This hydration procedure included placing a water soaked towel against the skin test site for one minute prior to actual spectra collection. A skin test site selected was an area of about eighty square centimeters, and about ten to twelve milligrams of each solution tested was applied to the skin test site area in the form of a thin film using a small paint brush. From the data collected, it was possible to calculate percentages of ingredients remaining on the skin following various soap wash sequences. The soap employed was a 0.5 weight percent solution of IVORY bar soap, and a soap rub is defined as two passes over the test area with the soap solution cupped in the palm of the hand. One soap wash procedure included fifteen soap rubs and ten rinse rubs under cool running tap water. The test site was the volar forearm. The test solutions were applied to the skin test site on the forearm in the form of a mixture of the various silicones and emollients, dissolved in a volatile silicone fluid of low viscosity, such as polydimethylcyclosiloxane which is a mixture of tetramer and pentamer having a viscosity of about 2.5 centistokes measured at 25°C. The solution contained five to ten percent by weight of the mixture in the solvent. The solvent was allowed to evaporate from the volar forearm region for fifteen to thirty minutes prior to the institution of the measurement procedures. The site was hydrated as noted above and initial spectrum was collected. The data included tests conducted with and without the presence of the various silicones in the test mixture, and at least two test runs were conducted for each mixture.

A simplified test procedure is illustrated as follows. A test area on the forearm was marked, and the test area was washed with the soap solution using fifteen rubs, followed by rinsing with ten rubs under cool running water. Excess moisture was blotted from the forearm with a towel. After one minute, the skin was hydrated for one minute using a towel saturated with water which was held loosely over the test area. Excess moisture was blotted, and at the end of thirty seconds a background scan was run. The test mixture was applied to the skin test area and the solvent allowed to evaporate. The skin was again hydrated for one minute and excess moisture was blotted off. After thirty seconds, a scan was run of the test area which represented an Initial Condition. The test area was washed with the soap solution using fifteen rubs followed by ten rinses, and the excess moisture was blotted off. After one minute, the skin was hydrated for one minute, blotted, and at the end of thirty seconds, a scan was run of the test area which represented a First

Soap Wash Condition. Similar steps were repeated for second, third, and fourth, soap wash conditions. Baselines for infrared bands were defined and band heights were measured. The percent ingredient remaining on the skin was calculated using this data.

The following tables set forth the results of the foregoing procedures, and illustrate the concept of the present invention of enhancing the durability of various emollients with certain functional silicones. The tables indicate that the functional silicones enhance emollient durability and therefore provide a viable solution to dry, chapped, and rough skin, which results when the emollients are removed by washing. The siloxanes are soap wash resistant and have shown minimal or no dermal irritation. The carboxyfunctional siloxane is known to possess the least dermal irritation of the siloxane types tested. The functional silicones used in the tables conform to the formula Me₃SiO(Me₂SiO)_x(MeRSiO)_ySiMe₃ in which Me is methyl and R is the functional group. Specifics of the R group and values of the integers x and y are set forth in the tables where appropriate. Unless otherwise indicated, the mixtures of emollient and silicone were in a ratio of four to one, and the mixtures were delivered in form of mixtures including a volatile cyclic siloxane. The compositions of the present invention may contain other adjuvants such as perfumes, fragrances, and preservatives, provided the addition of the adjuvant to the composition would not materially affect the basic and novel characteristics of the composition and would not materially change its fundamental characteristics.

TABLE I

EMOLLIENTS - NO SILICONE
PERCENT REMAINING

	Test Condition	Mink <u>Oil</u>	Lanolin <u>Oil</u>	Mineral <u>Oil</u>	<u>Petrolatum</u>
10	Initial	100	100	100	100
	lst wash	32	33	32	44
	2nd wash	19	21	16	34
15	3rd wash	14	15	. 6	19
	4th wash			4	16
20	5th wash			2	13

TABLE II

COMPOUNDS USED
AMINE AND AMIDE FUNCTIONAL SILICONES

30	Silicone	R-Group	<u>m%r</u>	<u>x</u>	Y	<u>x/y</u>
	A	iBuNH(CH ₂) ₂ NH ₂	1	97	1	91/1
	В	$iBuNH(CH_2)_2NH_2$	2	96	2	48/1
35	С	iBuNH(CH ₂)2NH ₂	5	188	10	19/1
	D	$iBuNH(CH_2)_2NH_2$	0.7	296	2	148/1
40	E	$iBuNH(CH_2)_2NH_2$	0.5	445.8	2.2	203/1
70	F	iBuNH(CH ₂) ₂ NH ₂	1.7	440.4	7.6	58/1
• .	G	$iBuNH(CH_2)_2NH_2$	0.25	796	2	398/1
45	Н	iBuNH(CH ₂) ₂ NHCOCH ₃	2	96	2	48/1

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TABLE III

PERCENT REMAINING MINERAL OIL AND SILICONES

	Test			<u>s</u> :	ILICON	<u>ES</u>			
	Condition	<u>H</u>	<u>A</u>	<u>B</u>	<u>C</u>	D	<u>E</u>	<u>F</u>	<u>G</u>
10	INITIAL	100	100	100	100	100	100	100	100
	lst wash	46	56	53	47	46	46	50	52
15	2nd wash	28	38	35	35	40	32	34	41
,,	3rd wash	22	32	28	30	32	32	28	32
	4th wash	20	30	24	27	26	26	21	30
20	5th wash	17	25	20	24	22	25	20	19

TABLE IV

PERCENT REMAINING MINERAL OIL AND SILICONES

30	Test <u>Condition</u>	Polydin 100DP	nethylsilo: 550DP	kane (1) 6800DP	<u>H</u>	Average <u>A-G</u>
	INITIAL	100	100	100	100	100
35	1st wash	35	35	45	46	50
	2nd wash	24	20	37	28	. 36
	3rd wash	20	18	31	22	31
40	4th wash	18	16	27	20	26
	5th wash	14		24	17	22

 $_{45}$ (1): 100 DP is 350 Centistoke fluid.

550 DP is hydroxy endblocked fluid with partial trimethyl capping.

6800 DP is a siloxane gum.

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TABLE V

% Mineral Oil Remaining

5	Sample <u>Number</u>	Polymer Description	Initial Condition	lst <u>Wash</u>	2nd <u>Wash</u>	3rd <u>Wash</u>	4th <u>Wash</u>	5th <u>Wash</u>
	1	No Silicone	100	32	16	6	4	2
10	2	н	100	56	34	25	20	14
	3	A	100	40	26	18	16	14
15	4	В	100	52	27	18	19	13
	5	С	100	40	28	25	18	14
	6	D	100	58	36	30	17	· 8
20	7	E	100	66	41	34	26	24
	8	F	100	58	37	28	17	15
25	9	G	100	33	30	18	20	5

TABLE VI

Compounds Used Carboxyfunctional Silicones

35	Reference	R-Group	<u> M%R</u>	<u>x</u>	Y	<u>x/y</u>
	J	iPr COOH	3	201.7	6.3	32/1
40	K	iPr COOH	0.7	296	2	148/1
40	L	iPr COOH	3.3	288	10	29/1
	М	iPr COOH	15	253	45	6/1

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TABLE VII

Percent Remaining Mineral Oil and Silicone

5		HIHELAT OIL AND SI	TICONE	
	Test <u>Condition</u>	<u>J</u>	<u>K</u>	<u>L</u>
10	INITIAL	100	100	100
	lst Wash	44	42	29
	2nd Wash	30	33	19
15	3rd Wash	24	28	14
	4th Wash	21	26	13
20	5th Wash	19	22	15

TABLE VIII

Mineral Oil and Silicone Percent Silicone Remaining

30	Test Condition	Polydi 100DP	methylsi 550DP	iloxane 6800DP	<u>H</u>	A-G <u>Amino</u> Avg.	J-K <u>Carboxylic</u> <u>Acid Avg.</u>
	INITIAL	100	100	100	100	100	100
35	lst Wash	35	35	45	46	50	43
55	2nd Wash	24	20	37	28	36	32
	3rd Wash	20	18	31	22	31	26
40	4th Wash	18	16	27	20	26	24
	5th Wash	14		24	17	22	20

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TABLE IX

Mineral Oil and Silicone % Mineral Oil Remaining

	Test Condition	No Sil.	<u>J</u>	<u>K</u>	<u>L</u>
10	INITIAL	100	100	100	100
	lst Wash	32	47	52	32
	2nd Wash	16	30	36	20
15	3rd Wash	. 6	19	24	13
	4th Wash	4	13	22	. 6
20	5th Wash	2	8	12	4

TABLE X

Two to One Mink Oil and Silicone

% Mink Oil Remaining

	Test	No	C	arboxylic Aci	.d
30	Condition	<u>Sil.</u>	<u>J</u>	<u>K</u>	<u>L</u>
	INITIAL	100	100	100	100
	lst Wash	32	46	45	30
35	2nd Wash	19	28	28	17
	3rd Wash	14	26	25	10

The mixture of emollient and silicone can be delivered to the skin in the form of emulsions, microemulsions, solutions, dispersions, lotions, gels, aerosols, solid sticks, ointments, and creams.

Claims

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A method of enhancing the durability of an emollient material on a substrate comprising forming a
mixture of an emollient material and an effective amount of an organosilicon compound, and applying
the mixture to the substrate to be treated, the organosilicon compound being selected from the groups
consisting of (a) aminofunctional polysiloxanes having the formula

 $R_{3-z}'Q_zSiO[R_2'SiO]_x[R'QSiO]_ySiQ_zR_{3-z}'$

wherein R' denotes an alkyl group of 1 to 4 carbons or a phenyl group, with the proviso that at least 50 percent of the total R' groups are methyl; Q denotes an amine functional substitutent of the formula —R"Z, wherein R" is a divalent alkylene radical of 3 to 6 carbon atoms or a radical of the formula —CH₂CH₂CH₂OCH₂-CHOHCH₂— and Z is a monovalent radical selected from the group consisting of —NR₂", —NR"(CH₂)_nNR₂"; and

wherein R''' denotes hydrogen or an alkyl group of 1 to 4 carbons, R'''' denotes an alkyl group of 1 to 4 carbons and n is a positive integer from 2 to 6; z has a value of 0 or 1; x has an average value of 25 to 3000; y has an average value of 0 to 100 when z is 1, and y has an average value of 1 to 100 when z is 0, (b) amidofunctional polysiloxanes having an average of 50 to 1000 siloxane units per molecule with an average of 1 to 50 of the siloxane units per molecule being amide-containing siloxane units bearing a substituent of the formula

wherein n is 0 or 1, R' denotes an alkylene radical of 3 to 6 carbon atoms, and R" denotes a hydrogen radical or an alkyl radical of 1 to 6 carbon atoms, X denotes an acyl radical of the formula

X' denotes a hydrogen radical or X, and R''' denotes an alkyl radical of 1 to 4 carbon atoms and substantially all other organic substituents in the polysiloxane being methyl groups, and (c) carboxyfunctional polysiloxanes having the formula

QMe₂SiO(Me₂SiO)_x(MeRSiO)_ySiMe₂Q

wherein Me is a methyl radical, R is a carboxyfunctional radical, said carboxyfunctional radical being selected from the group consisting of carboxyalkyl radicals and carboxythioalkyl radicals, Q is selected from the group consisting of R, Me and OH groups, x has a value of 1 to 1000, and y has a value of 1 to 100.

- 2. The method of claim 1 in which the mixture of the emollient and the polysiloxane includes a volatile cyclic siloxane solvent.
- 3. A skin conditioning composition of enhanced durability consisting essentially of a mixture of an emollient material and an effective amount of an organosilicon compound, the organosilicon compound being selected from the group consisting of (a) aminofunctional polysiloxanes having the formula
- 45 $R_{3-z}'Q_zSiO[R_2'SiO]_x[R'QSiO]_vSiQ_zR_{3-z}'$

wherein R' denotes an alkyl group of 1 to 4 carbons or a phenyl group, with the proviso that at least 50 percent of the total R' groups are methyl; Q denotes an amine functional substitutent of the formula —R"Z, wherein R" is a divalent alkylene radical of 3 to 6 carbon atoms or a radical of the formula —CH₂CH₂CH₂CCH₂-CHOHCH₂— and Z is a monovalent radical selected from the group consisting of —NR₂", —NR"(CH₂)_nNR₂"; and

wherein R" denotes hydrogen or an alkyl group of 1 to 4 carbons, R" denotes an alkyl group of 1 to 4

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carbons and n is a positive integer from 2 to 6; z has a value of 0 or 1; x has an average value of 25 to 3000; y has an average value of 0 to 100 when z is 1, and y has an average value of 1 to 100 when z is 0, (b) amidofunctional polysiloxanes having an average of 50 to 1000 siloxane units per molecule with an average of 1 to 50 of the siloxane units per molecule being amide-containing siloxane units bearing a substituent of the formula

wherein n is 0 or 1, R' denotes an alkylene radical of 3 to 6 carbon atoms, and R" denotes a hydrogen radical or an alkyl radical of 1 to 6 carbon atoms, X denotes an acyl radical of the formula

X' denotes a hydrogen radical or X, and R'" denotes an alkyl radical of 1 to 4 carbon atoms and substantially all other organic substituents in the polysiloxane being methyl groups, and (c) carboxyfunctional polysiloxanes having the formula

QMe₂SiO(Me₂SiO)_x(MeRSiO)_ySiMe₂Q

wherein Me is a methyl radical, R is a carboxyfunctional radical, said carboxyfunctional radical being selected from the group consisting of carboxyalkyl radicals and carboxythioalkyl radicals, Q is selected from the group consisting of R, Me and OH groups, x has a value of 1 to 1000, and y has a value of 1 to 100.

4. The composition of claim 3 in which the mixture of the emollient and the polysiloxane includes a volatile cyclic siloxane solvent.

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EUROPEAN SEARCH REPORT

EP 91 10 3231

	OCUMENTS CONSIDI	dication, where appropriate,	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CI.5)
tegory	of relevan	t passages		
×	EP-A-0 078 597 (TORAY SIL Page 6, paragraph 4 - page 1	ICONE CO., LTD) 11, paragraph 3, claims *	1-4	A 61 K 7/48 A 61 K 7/06
×	WO-A-8 904 161 (THE GILLE Page 2, line 22 - page 9, line	ETTE CO.) 28; examples; claims *	1,3	
X,D	US-A-4 559 227 (CHANDRA * Column 3, line 65 - column 8 column 11, line 13 *	et al.) 3, line 4; column 9, line 30	- 1,3	
×	EP-A-0 095 238 (DOW COR * Page 2, line 30 - page 7, line	NING CORP.)	1,3	
X	EP-A-0 174 097 (DOW COR Page 9, line 18 - page 10, line examples; claims	INING CORP.) ne 23; page 12, lines 9-31;	1,3	
×	US-A-4 501 619 (GEE) * Column 2, line 45 - column	10, line 16, claims *	1,3	
				TECHNICAL FIELDS SEARCHED (Int. CI.5)
				A 61 K
	The present search report has t			Evaminos
	Place of search	Date of completion of s	earch	Examiner
	The Hague	10 June 91		COUCKUYT P.J.R.
	CATEGORY OF CITED DOC X: particularly relevant if taken alone Y: particularly relevant if combined wi document of the same catagory		the filing date D: document cited L: document cited	for other reasons
	A: technological background O: non-written disclosure P: intermediate document T: theory or principle underlying the i		&: member of the s document	same patent family, corresponding